

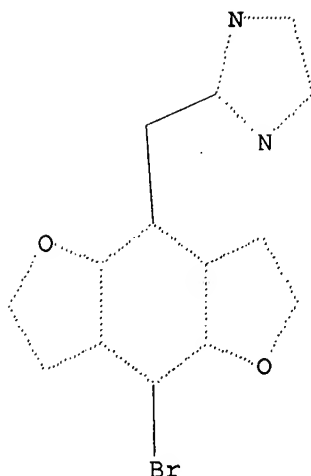
## EAST Search History

Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L1	494	548/311.1	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/01/04 17:48
L2	130	l1 and (ocular or eye)	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/01/04 17:52
L3	27	548/315.4 and (ocular or eye)	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/01/04 17:52
L4	10	548/347.1 and (ocular or eye)	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/01/04 17:53
L5	2	548/355.1 and (ocular or eye)	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/01/04 17:54
L6	31	549/429 and (ocular or eye)	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/01/04 17:55
L7	16	549/460 and (ocular or eye)	US-PGPUB; USPAT; EPO; JPO; DERWENT	OR	ON	2007/01/04 17:55

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548/211.1  
35.4  
347.1 355.1

549/429  
460



Structure attributes must be viewed using STN Express query preparation.

=> s l1 all

COMBINATION OF STRUCTURE AND TEXT TERMS NOT VALID

The query entered contains both search terms created by structure-building or screen commands and text search terms. L#s created via the STRUCTURE or SCREEN commands must be searched in the structures files separately from text terms or profiles. The L# answer sets from structure searches can be used in crossover searches and can be combined with text terms.

=> s l1 full

FULL SEARCH INITIATED 10:20:01 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 290 TO ITERATE

100.0% PROCESSED 290 ITERATIONS  
SEARCH TIME: 00.00.01

2 ANSWERS

L2 2 SEA SSS FUL L1

=> fil caplus

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

172.10

172.31

FILE 'CAPLUS' ENTERED AT 10:20:09 ON 04 JAN 2007

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FILE COVERS 1907 - 4 Jan 2007 VOL 146 ISS 2  
FILE LAST UPDATED: 3 Jan 2007 (20070103/ED)

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<http://www.cas.org/infopolicy.html>

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L3 1 L2

=> d ibib abs hitstr 1

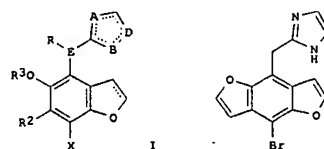
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L3 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2007 ACS on STN  
 ACCESSION NUMBER: 2003:511144 CAPLUS  
 DOCUMENT NUMBER: 139:85345  
 TITLE: Preparation of novel benzodifuranimidazolines and benzofuranimidazolines for the treatment of glaucoma  
 INVENTOR(S): Feng, Zixia; Hellberg, Mark R.  
 PATENT ASSIGNEE(S): Alcon, Inc., Switz.  
 SOURCE: PCT Int. Appl., 33 pp.  
 CODEN: PIXXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003053436	A1	20030703	WO 2002-US39316	20021209
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BE, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DS, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR				
TW 593302	B	20040621	TW 2002-91134883	20021129
CA 2469904	A1	20030703	CA 2002-2469904	20021209
AU 2002353088	A1	20030709	AU 2002-353088	20021209
EP 1455780	A1	20040915	EP 2002-790063	20021209
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK				
BR 2002015172	A	20041019	BR 2002-15172	20021209
CN 1606441	A	20050413	CN 2002-825464	20021209
JP 2005513103	T	20050512	JP 2003-554193	20021209
ZA 2004004473	A	20050607	ZA 2004-4473	20040607
US 2006009503	A1	20060112	US 2005-525410	20050128
PRIORITY APPLN. INFO.:				US 2001-343378P P 20011220
				WO 2002-US39316 W 20021209

OTHER SOURCE(S): MARPAT 139:85345  
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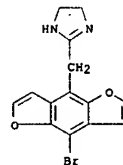
L3 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)



AB The title compds. [I; A, B, D = N, C, with the proviso that at least one of A, B or D = N; E = C, N; R = H, alkyl; R2, R3 = H, alkyl, alkenyl; or R2 and R3 together can form 5-6 membered ring; X = H, halo, alkyl, CF3], useful for lowering intraocular pressure and providing ocular neuroprotection, were prepared E.g., a multi-step synthesis of II.HCl, starting with bis(2-hydroxyethyl)hydroquinone, was given. The compound II.HCl showed IC50 of 0.46 nM and 6.4 nM against 5-HT2 and 5-HT1A receptor binding, resp. The compound II.HCl showed EC50 of 110 nM against  $\alpha$ 2A receptor binding. The pharmaceutical compns. comprising compds. I were claimed.

IT 554402-13-OP  
 RI: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)  
 (preparation of novel benzodifuranimidazolines and benzofuranimidazolines for the treatment of glaucoma)

RN 554402-13-0 CAPLUS  
 CN 1H-imidazole, 2-[(8-bromobenzo[1,2-b:4,5-b']difuran)methyl]-4,5-dihydro-, monohydrochloride (9CI) (CA INDEX NAME)



● HCl

L3 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)  
 REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE  
 FORMAT

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G2:H,CF2,CF3,X,[\*1],[\*2],[\*3],[\*4]

G3:H,[\*1],[\*2],[\*3],[\*4]

Match level :

1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:Atom 8:Atom 9:Atom 10:Atom  
11:Atom 12:Atom 13:Atom 14:Atom 15:Atom 16:CLASS 20:CLASS 21:CLASS 22:CLASS  
23:CLASS 24:CLASS 25:CLASS 26:CLASS 27:CLASS 28:CLASS 29:CLASS 34:CLASS  
36:CLASS

L4 STRUCTURE UPLOADED

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L4 HAS NO ANSWERS

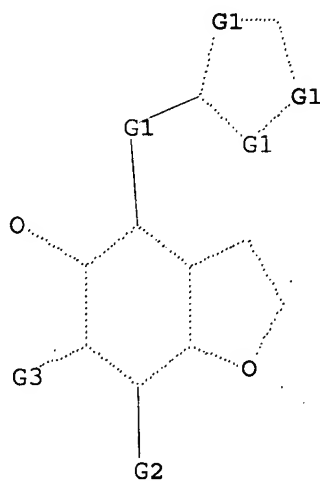
L4 STR

1

2

3

4



G1 C,N

G2 H,CF2,CF3,X,[@1],[@2],[@3],[@4]

G3 H,[@1],[@2],[@3],[@4]

Structure attributes must be viewed using STN Express query preparation.

=> s l4 full

FULL SEARCH INITIATED 10:28:36 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 184099 TO ITERATE

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100.0% PROCESSED 184099 ITERATIONS ( 1 INCOMPLETE) 9 ANSWERS  
SEARCH TIME: 00.00.01

L5 9 SEA SSS FUL L4

=> fil caplus  
COST IN U.S. DOLLARS

SINCE FILE	TOTAL
ENTRY	SESSION
172.10	350.15

FULL ESTIMATED COST

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE	TOTAL
ENTRY	SESSION
0.00	-0.78

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FILE COVERS 1907 - 4 Jan 2007 VOL 146 ISS 2  
FILE LAST UPDATED: 3 Jan 2007 (20070103/ED)

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=> s 15

L6 4 L5

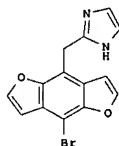
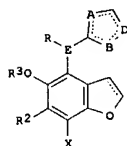
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10525410

L6 ANSWER 1 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN  
 ACCESSION NUMBER: 2003:51144 CAPLUS  
 DOCUMENT NUMBER: 139:85345  
 TITLE: Preparation of novel benzodifuranimidazolines and benzofuranimidazolines for the treatment of glaucoma  
 INVENTOR(S): Feng, Zixia; Heilberg, Mark R.  
 PATENT ASSIGNEE(S): Alcon, Inc., Switz.  
 SOURCE: PCT Int. Appl., 33 pp.  
 CODEN: FIKXD2  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003053436	A1	20030703	WO 2002-US39316	20021209
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR				
TW 593302	B	20040621	TW 2002-91134883	20021129
CA 2469904	A1	20030703	CA 2002-2469904	20021209
AU 2002353088	A1	20030709	AU 2002-353088	20021209
EP 1455780	A1	20040915	EP 2002-790063	20021209
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK				
BR 2002015172	A	20041019	BR 2002-15172	20021209
CN 1606441	A	20050413	CN 2002-825464	20021209
JP 2005513103	T	20050512	JP 2003-554193	20021209
ZA 2004004473	A	20050507	ZA 2004-4473	20040607
US 2006009503	A1	20060112	US 2005-525410	20050128
PRIORITY APPLN. INFO.:			US 2001-343378P	P 20011220
			WO 2002-US39316	W 20021209

OTHER SOURCE(S): MARPAT 139:85345  
 GI



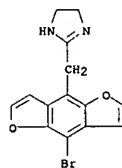
L6 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN  
 ACCESSION NUMBER: 2001:848926 CAPLUS  
 DOCUMENT NUMBER: 136:119162  
 TITLE: Preparation and characterization of a new solvent-free polymer electrolyte based on spiroketal structure  
 AUTHOR(S): Tsutsumi, Hiromori; Shirotani, Rumiko; Onimura, Kenjiro; Oishi, Tsutomu  
 CORPORATE SOURCE: Department of Applied Chemistry and Chemical Engineering, Faculty of Engineering, Yamaguchi University, Yamaguchi, 755-8611, Japan  
 SOURCE: Electrochemical and Solid-State Letters (2001), 4(12), A195-A196  
 CODEN: ESLEF6; ISSN: 1099-0062  
 PUBLISHER: Electrochemical Society  
 DOCUMENT TYPE: Journal  
 LANGUAGE: English  
 AB Solvent-free solid polymer electrolytes based on spiroketal structure were prepared and their properties were confirmed by conductance, differential scanning calorimetry, and X-ray diffraction measurements. The spiroketal polymer was synthesized from the bicyclic diketone and pentaerythritol. The spiro-polyketal (SP) dissolves lithium perchlorate and the conductivity of the (SP)1.5(LiClO4)1 complex is 4.24 x 10<sup>-5</sup> S cm<sup>-1</sup> at 30° and 3.83 x 10<sup>-4</sup> S cm<sup>-1</sup> at 60°.  
 IT 391671-11-7P  
 RL: POF (Polymer in formulation); PRP (Properties); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)  
 (Preparation and characterization of a new solvent-free polymer electrolyte based on spiroketal structure)  
 RN 391671-11-7 CAPLUS  
 CN Poly[3',6',6''a-diethyltetrahydrodispiro[1,3-dioxane-5,5'-(1,3)dioxane-2',2''(1''H)-pentalene]-2,5''(3''H)-diylidene] (9CI) (CA INDEX NAME)

\*\*\* STRUCTURE DIAGRAM IS NOT AVAILABLE \*\*\*  
 REFERENCE COUNT: 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE

FORMAT

L6 ANSWER 1 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)

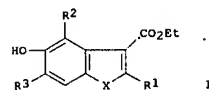
AB The title compds. [I; A, B, D = N, C, with the proviso that at least one of A, B or D = N; E = C, N; R = H, alkyl; R2, R3 = H, alkyl, alkenyl; or R2 and R3 together can form 5-6 membered ring; X = H, halo, alkyl, CF3], useful for lowering intraocular pressure and providing ocular neuroprotection, were prepared. E.g., a multi-step synthesis of II.HCl, starting with bis(2-hydroxyethyl)hydroquinone, was given. The compound II.HCl showed IC50 of 0.46 nM and 6.4 nM against 5-HT2 and 5-HT1A receptor binding, resp. The compound II.HCl showed EC50 of 110 nM against α2A receptor binding. The pharmaceutical compns. comprising compds. I were claimed.  
 IT 554402-13-0P  
 RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)  
 (Preparation of novel benzodifuranimidazolines and benzofuranimidazolines for the treatment of glaucoma)  
 RN 554402-13-0 CAPLUS  
 CN 1H-Imidazole, 2-[(8-bromobenzo[1,2-b:4,5-b']difuran)methyl]-4,5-dihydro-, monohydrochloride (9CI) (CA INDEX NAME)



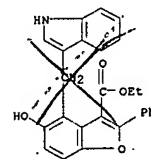
● HCl

REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE  
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L6 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN  
 ACCESSION NUMBER: 1995:796779 CAPLUS  
 DOCUMENT NUMBER: 124:29656  
 TITLE: Synthesis and some pharmacological properties of indole and benzofuran derivatives containing an imidazole pharmacophore  
 AUTHOR(S): Zotova, S. A.; Shvedov, V. I.  
 CORPORATE SOURCE: TsKhILS, VNIKhFI, Moscow, Russia  
 SOURCE: Khimiko-Farmatsevticheskii Zhurnal (1994), 28(2), 22-4  
 CODEN: KHFZAN; ISSN: 0023-1134  
 PUBLISHER: Meditsina  
 DOCUMENT TYPE: Journal  
 LANGUAGE: Russian  
 GI



AB Title compds. I (X = NMe, O; R1 = Me, Ph, CH2OPh, CH2SPh; R2 = imidazol-1-ylmethyl, Cl, H, CH2NMe2, indol-3-ylmethyl, CH2SCH2CONHPh; R3 = H, Br, imidazol-1-ylmethyl, CH2NMe2) were prepared by alkylation of imidazole with Mannich bases or by its reaction with bromoalkyl derivs. The synthesized compds. showed weak antimicrobial and anesthetic activity.  
 IT 171506-83-5P  
 RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)  
 (Preparation and pharmacol. activity of)  
 RN 171506-83-5 CAPLUS  
 CN 3-Benzofurancarboxylic acid, 5-hydroxy-4-(1H-indol-3-ylmethyl)-2-phenyl-, ethyl ester (9CI) (CA INDEX NAME)



10525410

L6 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1995:543579 CAPLUS

DOCUMENT NUMBER: 122:314550

TITLE: Preparation of (imidazolylalkyl)benzofurans and analogs as TXA2 synthetase and 5-lipoxygenase inhibitors and oxygen scavengers  
 INVENTOR(S): Ohuchida, Shuichi; Nambu, Fumio; Toda, Masaaki  
 PATENT ASSIGNEE(S): Ono Pharmaceutical Co., Ltd., Japan  
 SOURCE: Eur. Pat. Appl., 151 pp.  
 CODEN: EPXXDW  
 Patent

DOCUMENT TYPE:

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

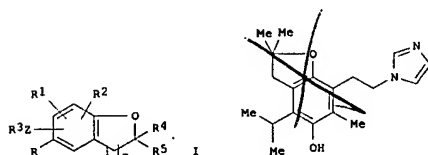
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 640609	A1	19950301	EP 1994-306175	19940822
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SE				
CA 2117551	A1	19950225	CA 1994-2117551	19940823
JP 07112980	A	19950502	JP 1994-221003	19940823
US 5534536	A	19960709	US 1994-294015	19940823
TW 403743	B	20000901	TW 1994-83107705	19940823
CN 1110969	A	19951101	CN 1994-117330	19940824
KR 192134	B1	19990615	KR 1994-20872	19940824
US 5750544	A	19980512	US 1996-635318	19960419
PRIORITY APPLN. INFO.:			JP 1993-231004	A 19930824
			US 1994-294015	A3 19940823

OTHER SOURCE(S):

MARPAT 122:314550

GI



AB Title compds. [I: R = OH, alkoxy, OBz, (di)(alkyl)amino, etc.; R1, R2 = H, halo, (cyclo)alkyl, alkoxy, etc.; R3 = 1 or 2 N-containing heterocyclyl;

R4, R5 = H, (phenyl)alkyl; CR4R5 = cycloalkyl; Z = alk(en)ylene, alkyleneoxy, (CH2)1-6OZ1; Z1 = 1,4-phenylene; n = 1-3] were prepared. Thus, title compound

II.HCl, prepared in 14 steps from 3-isopropyl-5-methylphenol, gave 74 and 92% inhibition of LTB4 and TXB2 production in whole human blood at 10 $\mu$ M in

L6 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN (Continued)

vitro.

IT 162962-70-1P

RL: BAC (Biological activity or effector, except adverse); BSU

(Biological

study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use);

BIOL (Biological study); PREP (Preparation); USES (Uses)

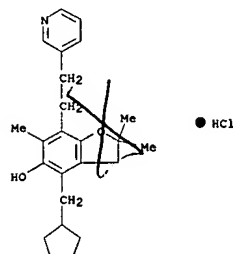
[preparation of (imidazolylalkyl)benzofurans and analogs as TXA2 synthetase

and 5-lipoxygenase inhibitors and oxygen scavengers)

RN 162962-70-1 CAPLUS

CN 5-Benzofuranol,

4-(cyclopentylmethyl)-2,3-dihydro-2,2,6-trimethyl-7-(2-(3-pyridinyl)ethyl)-, hydrochloride (9CI) (CA INDEX NAME)



IT 162963-70-4P 162963-71-5P 162963-72-6P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

[preparation of (imidazolylalkyl)benzofurans and analogs as TXA2 synthetase

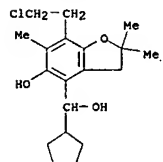
and 5-lipoxygenase inhibitors and oxygen scavengers)

RN 162963-70-4 CAPLUS

CN 4-Benzofuranmethanol, 7-(2-chloroethyl)- $\alpha$ -cyclopentyl-2,3-dihydro-5-hydroxy-2,2,6-trimethyl- (9CI) (CA INDEX NAME)

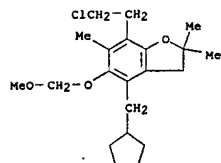
L6 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2007 ACS on STN

(Continued)



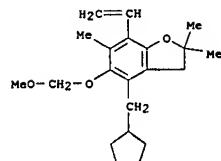
RN 162963-71-5 CAPLUS

CN Benzofuran, 7-(2-chloroethyl)-4-(cyclopentylmethyl)-2,3-dihydro-5-(methoxymethoxy)-2,2,6-trimethyl- (9CI) (CA INDEX NAME)



RN 162963-72-6 CAPLUS

CN Benzofuran, 4-(cyclopentylmethyl)-7-ethenyl-2,3-dihydro-5-(methoxymethoxy)-2,2,6-trimethyl- (9CI) (CA INDEX NAME)



Karen Cheng